We claim:

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1. A triazolopyrimidine of the formula I

$$\begin{array}{c|c}
N - N \\
N - N \\
N - R^2
\end{array}$$

5 in which the substituents are as defined below:

- R^1 is C_5 - C_{12} -alkyl or C_5 - C_{14} -alkoxyalkyl, where the aliphatic groups may be substituted by 1 to 3 of the following groups:
- 10 cyano, nitro, hydroxyl, C₃-C₆-cycloalkyl, C₁-C₆-alkylthio, and NR^aR^b;

 R^a , R^b are hydrogen or C_1 - C_6 -alkyl;

R² is CHR³CH₃, cyclopropyl, CH=CH₂ or CH₂CH=CH₂,

R³ is hydrogen, CH₃ or CH₂CH₃.

- 2. The compound of the formula I according to claim 1, in which R² is CHR³CH₃.
- 20 3. The compound of the formula I according to claim 1 or 2 in which R¹ is an unsubstituted straight-chain or mono-, di- or tribranched alkyl chain having up to 12 carbon atoms.
- 4. The compound of the formula I according to any of claims 1 to 3 in which R² is ethyl.
 - 5. The compound of the formula I according to any of claims 1 to 3 in which R² is isopropyl.
- 30 6. The compound of the formula I according to any of claims 1 to 3 in which R² is CHR³CH₃ and R³ is hydrogen, CH₂CH₃.
 - 7. 5-Ethyl-6-(1-methylheptyl)-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;

5-ethyl-6-octyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;

5-isopropyl-6-octyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;

5-ethyl-6-(3,5,5-trimethylhexyl)-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;

5-cyclopropyl-6-octyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;

5-ethyl-6-pentyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;

5-ethyl-6-hexyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;

5-ethyl-6-heptyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;

5-ethyl-6-nonyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;

5-ethyl-6-undecyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;

6-hexyl-5-isopropyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;

6-heptyl-5-isopropyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;

5-isopropyl-6-nonyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;

5-ethyl-6-(3-pentyloxypropyl)-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine.

8. A process for preparing compounds of the formula I according to any of claims 1
to 7 wherein ß-keto esters of the formula II,

$$RO$$
 R^{2}
 R^{1}
 R^{2}
 R^{2}

in which R is C₁-C₄-alkyl are reacted with 3-amino-1,2,4-triazole of the formula III

to give 7-hydroxytriazolopyrimidines of the formula IV

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which are halogenated to give compounds of the formula V

$$\begin{array}{c|c}
 & \text{Hal} \\
 & \text{N} \\
 & \text{N} \\
 & \text{N} \\
 & \text{R}^2
\end{array}$$

in which Hal is chlorine or bromine and V is reacted with ammonia.

- 20 9. A compound of the formula IV or V as set forth in claim 8.
 - A process for preparing compounds of the formula I according to any of claims 1 to 7 wherein acyl cyanides of the formula VI,

$$\begin{array}{ccc}
 & & & VI \\
 & & & & VI \\
 & & & & & & VI
\end{array}$$

- are reacted with 3-amino-1,2,4-triazole of the formula III as set forth in claim 8.
 - 11. A fungicidal composition comprising a solid or liquid carrier and a compound of the formula I according to any of claims 1 to 7.

- 12. Seed comprising a compound of the formula I according to any of claims 1 to 7 in an amount of 1 to 1000 g per 100 kg.
- A method for controlling phytopathogenic harmful fungi, wherein the fungi or the
 materials, plants, the soil or seed to be protected against fungal attack are
 treated with an effective amount of the compound of the formula I according to
 any of claims 1 to 7.